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ORIGINAL RESEARCH

Short running header: CDK4/6 and mTOR inhibitors synergistically limit growth of DIPG cells.

Daniel J. Asby et al

Combined use of CDK4/6 and mTOR inhibitors induce synergistic growth arrest of diffuse intrinsic pontine glioma cells via mutual downregulation of mTORC1 activity

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Abstract:

Background: Diffuse intrinsic pontine glioma (DIPG) is a lethal type of pediatric brain tumor that is resistant to conventional chemotherapies. Palbociclib is a putative novel DIPG treatment that restricts the proliferation of rapidly dividing cancer cells via selective inhibition of cyclin-dependent kinase (CDK) 4 and CDK6. However, implementing palbociclib as a monotherapy for DIPG is unfeasible, as CDK4/6 inhibitor resistance is commonplace and palbociclib does not readily cross the blood-brain barrier (BBB) or persist in the CNS. We aimed to use palbociclib to inhibit growth of DIPG cells, in combination with the rapamycin analogue temsirolimus, which is known to ameliorate resistance to CDK4/6 inhibitors and inhibit BBB efflux.

Materials and methods: We tested palbociclib and temsirolimus in three patient-derived DIPG cell lines. The expression profiles of key proteins in the CDK4/6 and mammalian target of rapamycin (mTOR) signaling pathways were assessed, respectively, to determine feasibility against DIPG. We also investigated effects on cell viability and examined *in vivo* drug toxicity.

Results: Immunoblot analyses revealed palbociclib and temsirolimus inhibited CDK4/6 and mTOR signaling, through canonical perturbation of phosphorylation of the retinoblastoma (RB) and mTOR proteins, respectively, while we also observed non-canonical downregulation of mTOR by palbociclib. We demonstrated that palbociclib and temsirolimus inhibited cell proliferation in all three DIPG cell lines, acting synergistically in combination to further restrict cell growth. Flow cytometric analyses revealed both drugs caused G₁ cell cycle arrest and clonogenic assays showed non-reversible effects on cell proliferation. Palbociclib did not elicit neurotoxicity in primary cultures of normal rat hippocampi or when infused into rat brains.

Conclusion: These data illustrate the *in vitro* antiproliferative effects of CDK4/6 and mTOR inhibitors in DIPG cells. Direct infusion of palbociclib into the brain, in combination with systemic delivery of temsirolimus, represents a promising new approach to developing a much-needed treatment for DIPG.

Keywords: Palbociclib, temsirolimus, brain tumor, DIPG, retinoblastoma protein, cyclin-dependent kinase

Introduction

Diffuse intrinsic pontine glioma (DIPG) is a lethal high-grade pediatric glioma that accounts for up to 85% of all brainstem gliomas.¹ Approximately 100-150 new cases of DIPG in children are recorded each year in the USA.² The genetic basis of DIPG is complex and several genetic and epigenetic alterations are associated with the disease. However, approximately 80% of DIPG tumors exhibit the missense mutation lysine 27 to methionine (K27M) in the genes that encode histones H3.1 and H3.3 (H3.1K27M and H3.3K27M, respectively).³ DIPG is difficult to treat due to the heterogeneity generated from the various mutations associated with the disease, as well as its sensitive location in the brainstem. Fractionated radiation is the conventional treatment but has no long-term benefit, with the two-year survival rate currently below 20%.¹ Chemotherapies have also proved ineffective and new treatments are urgently required. The results of combinatorial approaches utilizing synergistic targeted therapies have been encouraging.³

Palbociclib (also known as PD-0332991) is a selective inhibitor of cyclin-dependent kinases (CDK) 4 and 6.⁴ Both CDK4 and 6 form functionally identical heterodimeric complexes with cyclin D1 (cycD1), cycD2 or cycD3, which phosphorylate and inactivate retinoblastoma protein (RB).⁵ Inactivation of RB relieves negative regulation of the E2F transcription factor, which facilitates progression through the G₁/S transition in the cell cycle, permitting DNA synthesis and cell proliferation. Palbociclib inhibits the kinase activity of CDK4/6, thereby preventing phosphorylation of RB and cell cycle progression. This leads to cell cycle arrest in the G₁ phase of the cell cycle. Thus, treatment with palbociclib results in cells accumulating in G₁ and inhibition of cell proliferation. However, malignancies can develop resistance to CDK4/6 inhibitors,⁶ putatively via cycD1 expression flux, with down-regulation of the mammalian target of rapamycin (mTOR) representing one approach to abrogate this.⁴

Temsirolimus (previously known as CCI-779) is an ester analogue, or rapalogue, of the mTOR inhibitor rapamycin, which possesses increased aqueous solubility and improved pharmacokinetics compared to its parent molecule.⁷ Temsirolimus is antitumorigenic *in vitro* through targeted inhibition of the mTORC1 complex and was approved for intravenous systemic delivery for the treatment of renal cell carcinoma in 2007.⁸ The effectiveness of temsirolimus as a single agent therapy against other types of cancer has been limited,⁹ though it has shown some capacity to treat glioblastoma¹⁰ and has been shown to readily cross the blood-brain barrier (BBB).¹¹ Temsirolimus

combination therapies are currently being trialed for treatment of central nervous system (CNS) tumors, where it has been found to be well tolerated by patients and appears amenable to combination with other targeted therapies.¹²

We used three cell lines encompassing both archetypal DIPG histone mutations (H3.1K27M found in SU-DIPG IV cells and H3.3K27M found in SF7761 and SF8628 cells) to demonstrate that by selectively disrupting the CDK4/6-cycD1-RB and mTOR signaling pathways, sustained inhibition of DIPG cell proliferation can be achieved through synergistic cytostatic-driven effects on cell growth, increasing the therapeutic potential of these two candidate DIPG therapeutics.

Material and methods

Reagents were purchased from ThermoFisher Scientific (Loughborough, UK) unless otherwise stated. Palbociclib and temsirolimus were purchased from Selleckchem (Houston, Texas, USA). Drug synergy analysis, based on cell viability assays, compared single-agent treatments to equivalent combination treatments, as previously described.¹³ Statistical analyses were performed using GraphPad Prism 6 and MiniTab 17. All values are expressed as mean of triplicate determinations \pm SEM. Values of $p < 0.05$ were considered statistically significant.

Cell culture and cell treatments

Patient-derived SF7761 and SF8628 cell lines were isolated from DIPG tumor tissue acquired by the University of California San Francisco (UCSF) Tissue Bank. SU-DIPG IV cells were isolated from a DIPG patient at Stanford University. All procedures were conducted with Institutional Review Board approval. SF7761 and SF8628 cells were obtained from Nalin Gupta (UCSF) and SU-DIPG IV from Michelle Monje (Stanford University) via Material Transfer Agreements. Cells were authenticated by short tandem repeat (STR) profiling (Public Health England, UK). Cells were used within 10 passages from thawing and confirmed to be mycoplasma free (in-house testing). SF7761 and SF8628 culture has been described previously.¹³ SU-DIPG IV cells were grown in tumor stem media: Dulbecco's modified Eagle medium / Ham's F-12 (DMEM/F12) and Neurobasal-A media [1:1 ratio], with B27 neural cell culture supplement (2%), human basic fibroblast growth factor (hFGF-basic; 20 ng/ml; Peprotech, London, UK), mouse epidermal growth factor (mEGF; 20 ng/ml; Peprotech), human

platelet-derived growth factor AA (hPDGF-AA; 10 ng/ml; Generon, Maidenhead, UK), hPDGF-BB (10 ng/ml; Generon) and heparin (2 mg/ml, StemCell Technologies, Grenoble, France). Cells were seeded 16 hours prior to treatment in all instances and maintained at 5% CO₂ and 37°C. Cells were treated with drugs for 24 hours unless stated otherwise. Serially diluted stock solutions of palbociclib and temsirolimus were reconstituted in artificial cerebrospinal fluid (Torbay Pharmaceuticals, Paignton, Devon, UK) and dimethyl sulfoxide, respectively.

Protein immunoblotting

Immunoblotting was performed as previously described.¹⁴ Nitrocellulose membranes were probed overnight at 4°C with the following antibodies at their respective dilutions. From Cell Signaling Technology (Danvers, Massachusetts, USA): RB (4H1) #9309 1:1000; phospho-RB (Ser807/811) #8516 1:1000; CDK6 (D4S8S) #13331 1:500; cyclin D1 (92G2) #2978 1:500; mTOR #2972 1:1000; phospho-mTOR (Ser2448) #2971 1:1000; S6 kinase (49D7) #2708 1:1000; phospho-S6 kinase (Thr389, 108D2) #9234 1:1000; 4EBP1 (53H11) #9644 1:2000; phospho-4EBP1 (Thr37/46) #9459 1:500; Rictor (53A4) #2114 1:1000; phospho-rictor (Thr1135, D30A3) #3806 1:1000; anti-rabbit HRP #7074 1:2000; and anti-mouse HRP #7076 1:1000. In addition to p16^{INK4A} #MAB4133 (Millipore, Watford, UK) and CDK4 #559677 (Becton Dickinson, Wokingham, UK). Densitometry values are the mean of triplicate determinations and representative blots are shown.

Cell viability assays

Cell viability and cytotoxicity were measured with a two-color fluorescent-based live/dead kit (Life Technologies, Paisley, UK) utilizing calcein acetoxymethyl (calcein-AM) and ethidium homodimer-1 (ethD-1). Cells were seeded in 96-well plates and incubated for 16 hours prior to drug treatments. Cells were dosed with various concentrations of each drug, as described in the Results section below. Cells were then washed with phosphate-buffered saline and incubated with 2 µM calcein-AM and 4 µM ethD-1 for 30 minutes. Fluorescence was measured on a microplate reader (FLUOstar Omega; BMG Labtech, Aylesbury, UK) and normalized against appropriate controls.

Flow cytometry

Cells were seeded on 12-well plates and dosed with palbociclib, temsirolimus, or both, as described. SF7761, SF8628 and SU-DIPG IV cells were seeded at 3.5×10^5 , 0.4×10^5 and 1×10^5 cells per well, respectively, for experiments requiring 24 hours drug treatments; and at 1×10^5 , 0.2×10^5 and 0.4×10^5 cells per well, respectively, for 72 hours experiments. Cells were then detached from the growing surface using Accutase solution and collected. For cell viability analysis, calcein-AM and ethD-1 were diluted in PBS at 1:10,000 and 1:250, respectively, prior to addition to cell suspensions at a 1:1 ratio. For cell cycle analyses, DRAQ5 (eBioscience, San Diego, California, USA) was diluted 1:200 in PBS and added to cell suspensions at a 1:1 ratio.

Clonogenic assays

SU-DIPG IV cells were seeded at 800 cells per 6-cm dish, 16 hours prior to treatments described in the text. Cells were processed as previously described.¹⁴ At the end of the experiment cells were fixed by addition of methanol/acetic acid (3:1) for 5 minutes, then stained with 0.5% (v/v) crystal violet solution (diluted in methanol) for 10 minutes. Colonies were counted using the colony area plugin for ImageJ (version 1.46r; National Institutes of Health, USA).

In vitro neurotoxicity assessment via fluorescent

immunocytochemistry

Experiments were performed on rat E18 hippocampal cultures grown on poly-D-lysine coated glass coverslips after cell extraction had been performed using previously described protocols.¹⁵ Palbociclib toxicity was assessed after 24 hours of treatment using immunocytochemistry assays performed on untreated and palbociclib treated primary hippocampal cells as described previously.¹³

In vivo neurotoxicity assessment via fluorescent

immunohistochemistry

Targeted delivery of palbociclib to the brain striatum of Wistar rats was achieved using a custom cannula system and rate-controlled microinfusion pump, as previously described.¹⁵ The

animals used in this study were handled according to the protocols approved by the ethical committee of University of Bristol and all the protocols were performed in accordance with the UK Animal Scientific Procedures Act 1986. Immunohistochemical analysis of tissue sections was performed as previously described.¹⁵ Briefly, animals were transcardially perfused with 4% paraformaldehyde and the brains removed, then fixed using 4% paraformaldehyde. Rat brains were then cut into 35 μ m thick sections at -20°C and mounted on microscope slides. Anti-NeuN (1:100; Millipore, Burlington, MA, USA) and anti-GFAP (1:300; Millipore) primary antibodies were used together with standard immunofluorescent protocols to identify any neuronal disruption and gliosis, respectively.

Results

Combination treatment with palbociclib and temsirolimus inhibit both the CDK4/6-cycD1-RB and mTORC1 signaling pathways in DIPG cells

Barton et al, 2013 previously showed that palbociclib was effective at inhibiting the CDK4/6-cycD1-RB signaling pathway at sub-micromolar concentrations in genetically engineered murine DIPG cells.¹⁶ Furthermore, analogues of temsirolimus, such as everolimus, have been shown to be effective at inhibiting cell viability at low-micromolar concentrations in non-DIPG brain malignancies, such as glioblastoma.^{17,18} Thus, in order to establish whether palbociclib and temsirolimus could be clinically relevant to DIPG, we initially examined their effects on the expression of key proteins in the CDK4/6-cycD1-RB and mTOR signaling pathways in three *ex vivo* human DIPG cell lines, at concentrations known to elicit strong on-target effects (0.5 μ M and 10 μ M, respectively).

Heterodimeric CDK4/6-cycD complexes phosphorylate, and consequently, inhibit RB activity. Therefore, hypophosphorylated RB is an indicator of inhibition of CDK4/6 and release of the block on RB activity. We found palbociclib effectively inhibited CDK4/6 after 24 hours, eliciting marked decreases in the phosphorylation of RB in all three cell lines (Fig. 1), and dose-dependently across a range of doses in SF7761 (Fig. 2). As palbociclib is thought to have maximal activity in situations where RB is intact and p16^{INK4A} is lacking, we wished to clarify that p16^{INK4A} expression was absent in

our cells (although deletion of CDKN2A and a lack of p16^{INK4A} protein expression has been shown previously in SF7761,¹⁹ SF8628²⁰ and SU-DIPG IV²⁰). As our cell lines all lacked p16^{INK4A} expression (Fig. S1), we suggest that in DIPG cells palbociclib acts as a proxy for the p16^{INK4A}-mediated inhibition of CDK4/6, and the consequential attenuation of cell cycle progression, normally found in non-cancerous cells.⁴ Furthermore, as the expression of CDK4, CDK6 and cycD1 were largely unaffected by palbociclib treatment, it appears more likely that the observed impact of palbociclib on cell proliferation resulted from its archetypal effects on CDK4/6 inhibition alone, rather than any off-target effects or non-specific hypertoxicity affecting the CDK4/6-cycD-RB pathway.

In contrast, temsirolimus had a limited effect on RB activation in the cell lines tested (Fig. 1), and only demonstrated an effect on RB at the highest doses in SF7761 (i.e. 10-40 μ M; Fig. 2). Concurrent use of both drugs potentiated inhibition of CDK4/6 by palbociclib and enhanced RB activation, indicative of antiproliferative effects. Immunoblotting also revealed that the combination treatment produced a conspicuous decrease in CDK expression, which was not readily observable following single-agent treatments. This may also contribute to inhibition of cell proliferation.

Temsirolimus is a rapamycin analogue and hence a specific inhibitor of mTOR. It binds with FK506-binding protein 12 (FKBP12), which then directly binds mTORC1 and obscures the correct alignment of substrates to its catalytic cleft; preferentially inhibiting phosphorylation of the mTOR substrate p70 ribosomal protein S6 kinase (S6K) over 4E-binding protein 1 (4EBP1).²¹ In all three cell lines, temsirolimus elicited a significant decrease in mTORC1 activity, indicated by decreased phosphorylation of mTOR and its downstream targets S6K and 4EBP1 (Fig. 1). This effect was dose-dependent in SF7761 and no phosphorylated mTOR protein was detectable following treatment with 20 μ M temsirolimus (Fig. 2). Dose-dependent reduction of cycD1 also resulted from temsirolimus treatment, which showed some correlation with reduced mTOR activity in the lower dose range (fold change at: 0.2 μ M = 0.65 and 0.48; 2 μ M = 0.38 and 0.21; 10 μ M = 0.33 and 0.08; 20 μ M = 0.17 and 0.01; 40 μ M = 0.07 and 0.01 μ M; for cycD1 and p-mTOR/mTOR, respectively). This reflects the role of mTORC1 as a positive regulator of the translation of cycD1 protein,²² indicating on-target effects of temsirolimus in DIPG cells.

Furthermore, palbociclib potentiated inhibition of mTORC1 by temsirolimus, with the addition of 0.5 μ M palbociclib bringing about additional significant down-regulation of mTOR, S6K and 4EBP1 phosphorylation. Palbociclib's augmentation of the effectiveness of temsirolimus on mTOR pathway

inhibition shows hallmarks of being more than just an additive effect, because when palbociclib was used as a single drug at this low dose it had an unremarkable effect on mTOR, S6K or 4EBP1. With single agent temsirolimus eliciting substantial reductions in mTOR activity, palbociclib's potentiation of the direct effect of temsirolimus on mTOR was difficult to quantify, with increased effects on mTOR slightly observable in only one cell line (SF7761; Fig. 2). However, increased effects on mTOR following combination treatment were observed indirectly via increased hypophosphorylation of 4EBP1 in all three cell lines (Fig. 1). For example, in SF7761, using 0.02 μ M temsirolimus in combination with 0.5 μ M palbociclib led to a fold change of 0.32 in mTOR phosphorylation, whereas the same dose of temsirolimus only elicited a fold change of 0.93 for the single agent treatment (Fig. 1). Likewise, S6K and 4EBP1 showed significantly increased hypophosphorylation in SF7761 across a range of doses following combination treatment, as compared to temsirolimus alone (e.g. p-S6k/S6K: fold change at 0.02 μ M temsirolimus = 0.90 and 0.01 for temsirolimus and temsirolimus + palbociclib, respectively; and p-4EBP/4EBP: fold change at 0.02 μ M temsirolimus = 0.97 and 0.53 for temsirolimus and temsirolimus + palbociclib, respectively).

Concurrent use of temsirolimus with palbociclib also resulted in increased activation of RB, despite the equivalent single agent doses having minimal effects on RB phosphorylation (Fig. 1). Specifically, in SF7761 cells, 0.5 μ M PD and 0.02 μ M TM only elicited a fold-change in RB phosphorylation of 0.5 and 0.93, respectively. In combination, these doses caused a fold-change of 0.06. Furthermore, combination treatment produced a conspicuous decrease in CDK expression, which was not readily observed following single agent treatments (Fig. 1). Finally, use of both drugs together promoted hypophosphorylation of the mTORC2-associated rictor protein (Fig. 2), signifying rictor activation and enhanced mTORC2 activity. As S6K is a negative regulator of rictor,²³ it is unsurprising that the increased inhibition of S6K observed following combination treatment, stimulated rictor activation. However, this is an undesirable side-effect of an antitumorigenic treatment because increased mTORC2 activity signifies increased cell growth and proliferation. Nevertheless, considering the increased antiproliferative effects of the drugs in combination observed in our other assays (see below), we propose the concurrent use of both drugs establishes a cellular environment not conducive to cell expansion, thus mTORC2 activation is ineffectual.

CDK4/6 and mTOR inhibition decrease viability in DIPG cells without triggering extensive cytotoxicity

Once we had established that palbociclib and temsirolimus had significant on-target effects on DIPG cells, we attempted to establish whether these molecular effects translated to significant inhibition of DIPG cell growth. We first measured changes in cell viability in our three DIPG cell lines treated with a range of concentrations of either palbociclib or temsirolimus (200 nM - 200 μ M). Following 24 hours treatment with palbociclib, cell viability was found to decrease in a dose-dependent manner in all three cell lines (Fig. 3A). A highly significant reduction in viability was reached by 15 μ M palbociclib in all cell lines. However, significant cytotoxicity was only evident in cells at 25 μ M palbociclib (Fig. 3B). This implies a cytostatic mode of action at IC₅₀ doses of palbociclib (Table 1 and Fig. S2). Similarly, temsirolimus also provoked a dose-dependent loss in cell viability in all three cell lines (Fig. 3C) and showed cytostatic traits. Specifically, temsirolimus elicited significant loss of viability in SF7761 and SU-DIPG IV following treatment with 10 μ M temsirolimus, whilst only causing substantial toxicity at 25 μ M (Fig. 3D); this is higher than the calculated IC₅₀ values of 20.9 and 16.8 μ M, for SF7761 and SU-DIPG IV, respectively (Table 1 and Fig. S2). In SF8628, no significant cytotoxicity following temsirolimus treatment was observed at any of the concentrations tested (Fig. 3D).

Palbociclib and temsirolimus act synergistically to restrict the growth of DIPG cells

It has been shown previously in other malignancies that overcoming CDK4/6 inhibitor resistance and increasing the therapeutic potency of CDK4/6 inhibitors can be achieved by combining this class of therapeutic with other pathway selective agents.²⁴ As palbociclib and temsirolimus appeared effective at inhibiting DIPG cell growth as monotherapies, we next assessed the effect of the drugs in combination on cell viability. To achieve this, we treated cells with a variable concentration of temsirolimus (200 nM-200 μ M) in combination with a single fixed dose of palbociclib for 24 hours (either 0, 2, 10, 12, 15 or 25 μ M). Combination treatments demonstrated that the two

drugs worked effectively in combination and increasing the concentration of the fixed dose of palbociclib triggered greater reductions in cell viability (Fig. 4). Quantitative assessment of synergy generated combination indices that were indicative of additive or synergistic effects. For example, in SF7761 and SF8628 cells a 15 μ M fixed dose of palbociclib yielded highly synergistic combinations with various doses of temsirolimus, exclusively producing combination index values <1 , most lying between 0.3 and 0.7, consistent with 'true synergism'.²⁵ In experiments involving SU-DIPG IV, a lower dose of 10 μ M palbociclib was sufficient to bring about strong synergistic effects on cell viability.

A more detailed examination of cell viability using flow cytometry (Fig. 5) supported our initial findings, whilst also indicating that although palbociclib caused only limited cell death at lower doses, cell numbers generally decreased steadily; below the percentage of live cells in all instances (Fig. 5, left panels). Doses of palbociclib above 25 μ M appeared to be highly toxic, causing considerable cell death, consistent with our initial cytotoxicity assays (Fig. 3B). Thus, palbociclib doses of this magnitude appeared fundamentally cytotoxic and were excluded from the experiments that followed. Cytostatic-driven reductions in cell viability were more evident in cells treated with temsirolimus. In SF7761 and SF8628 temsirolimus produced negligible cell death at all concentrations up to 50 μ M, whilst cell numbers consistently decreased in a dose-dependent manner relative to untreated cells (Fig. 5, center panels). Coadministration of a low fixed dose of palbociclib (either 15 μ M for SF7761 and SF8628, or 10 μ M for SU-DIPG IV cells), combined with varying concentrations of temsirolimus potentiated the effects on cells (Fig. 5, right panels), provoking a significant decrease in the number of cells, together with cell viability (particularly in SF7761 and SU-DIPG IV). Implying a shift from a cytostatic to cytotoxic mode of action, indicative of increased anti-tumorigenic properties. These doses of palbociclib were chosen as they were close to the drug's IC₅₀ and also elicited strong synergism with temsirolimus (established in the previous experiments above), while they were also largely sub-lethal in the single drug flow cytometric analyses.

Combined CDK4/6 and mTOR inhibition cytostatically inhibit proliferation and trigger G₀-G₁ cell cycle arrest

Having observed that the combination of palbociclib and temsirolimus potentiated each drug's cytostatic effects, we next investigated whether these effects continued beyond the initial 24-hour

time-period. Low, sub-lethal doses of each drug were selected based on the previous data, as well as cell line-specific drug sensitivity and some preliminary experiments examining the effects of the drugs over time (data not shown). These criteria led to us utilizing a dose of 2 μ M palbociclib for all three cell lines, as this drug concentration had no effect on cell death in any of the three cell lines, and only a very limited effect on cell proliferation in SF8628 and SU-DIPG IV. A concentration of 10 μ M temsirolimus was utilized for SF7761 and SF8628, while a lower dose of 0.2 μ M temsirolimus was used for SU-DIPG IV, as this cell line was more sensitive than the other cell lines to temsirolimus, and showed significant loss of viability at 10 μ M in the previous flow cytometric analyses. Throughout the 72 hours viability in control cells remained high (Fig. 6A-C, left). Similarly, treatment with either palbociclib or temsirolimus did not impact on cell viability. However, cells treated with palbociclib or temsirolimus clearly demonstrated diminished proliferation, which intensified over time (Fig. 6A-C, center and right, respectively).

Effects on cell proliferation were investigated further via cell cycle analyses. In SF7761 cells, treatment with palbociclib and temsirolimus for 24 hours caused a marked increase in the percentage of cells in the G₀-G₁ phase of the cell cycle, consistent with CDK4/6 inhibition and mTOR inhibition, respectively (Fig. 7A). G₀-G₁ arrest persisted over 72 hours in all three cell lines tested, with SU-DIPG IV demonstrating the greatest amount of cell cycle arrest (Fig. 7B, 7C and S3). To further assess the antiproliferative effects of the drugs, we performed clonogenic assays using SU-DIPG IV cells, which unlike SF7761 and SF8628, exhibit strong intrinsic clonogenicity. We found palbociclib and temsirolimus inhibited colony formation time-dependently as single agents, and concurrent treatment with both drugs potentiated the effect of temsirolimus at all three time points (Fig. 7D and S4). The clonogenicity of SU-DIPG IV showed very high sensitivity to palbociclib in our preliminary combination experiments, and thus a dose of 0.5 μ M palbociclib (in line with our initial immunoblotting assays) was used in combination with temsirolimus, which was sufficient to considerably augment the ability of temsirolimus to reduce colony numbers. Taken together these data suggest the drugs provided sustained antiproliferative effects, via cell cycle arrest.

Palbociclib does not cause significant neurotoxicity in normal brain tissue

It has previously been demonstrated that the lipophilic nature of temsirolimus enables it to readily cross the blood-brain barrier (BBB).¹¹ In contrast, palbociclib does not readily cross the BBB.²⁶ To address this issue, we propose it would be prudent to deliver palbociclib directly into the brain of DIPG patients to circumvent the BBB, while temsirolimus could be infused intravenously, which does not pose any known toxicity issues.¹² With this in mind, we examined the effect of palbociclib on normal brain tissue. To achieve this, we initially exposed normal neurons and glial cells within primary rat hippocampal cultures to 2 μ M palbociclib and observed no discernable toxic effects, relative to the general morphology and numbers of cells seen in vehicle-treated cultures (Fig. 8). We went on to investigate the toxicity of palbociclib *in vivo* by carrying out direct infusions of either 100 μ M or 400 μ M palbociclib into the striatum of normal rat brains. Analyses of the brain tissue showed no significant neurotoxicity caused by palbociclib, as compared to control animals (Fig. 9).

Discussion

DIPG is a highly heterogeneous disease characterized by lethal tumors that differ in cellular origin and pathogenesis. In part, this heterogeneity is derived from the genetic basis of the disease and two archetypal causative mutations.^{27,28} In the present study we have shown that palbociclib and temsirolimus can reduce DIPG cell proliferation *in vitro* in three different cell lines encompassing the two histone H3 mutations most frequently associated with DIPG (H3.3K27M represented by SF7761 and SF8628; and H3.1K27M in SU-DIPG IV). On the whole we found that palbociclib and temsirolimus was more efficacious in SU-DIPG IV cells; suggesting that DIPG tumors harboring the H3.1K27M mutation may be more suited to the proposed drug combination. Though SF7761 and SF8628 cells both harbor the H3.3K27M mutation, SF7761 was generally more sensitive to drug treatments. This variation in the two H3.3K27M cell lines likely relates to other genotypic differences, and phenotypic differences, that exist between these cells. Crucially, the SF7761 cell line has been modified to express telomerase, whereas SF8628 is unmodified and has been established directly from DIPG biopsy. Consequently, SF7761 cells can potentially survive indefinitely *in vitro*, whereas

SF8628 cells senesce over time. Furthermore, SF7761 have a shorter doubling time compared to SF8628 (Fig. 6), are typically much smaller cells and are cultured in a different culture media formulation. These cell lines have also been shown to have distinctive endogenous genetic profiles beyond the shared H3 histone mutation. For example, expression of the nuclear kinase *WEE1* is much higher in SF8628 compared to SF7761.²⁹ Asymmetrical expression of this important cell cycle regulator could conceivably contribute to the dissimilar response to our proposed novel drug combination, which specifically targets cell cycle arrest. Irrespective of these differences, the combination of palbociclib and temsirolimus brought about strong growth inhibition in both SF7761 and SF8628.

While many novel DIPG therapeutics have been proposed in recent years, none have been successfully implemented.^{30, 31} Single-agent rapalogue treatments for CNS tumors, including DIPG, have had limited success.³²⁻³⁶ However, temsirolimus has been well tolerated in pediatric patients.¹² The situation is similar for CDK4/6 inhibitors, where use of these drugs as robust single-agent treatments for different cancers has proven unconvincing.^{32, 38-40} Nevertheless palbociclib has been well tolerated, increased survival in a DIPG mouse model¹⁶ and can elicit anti-tumor immunological effects.⁴¹ Both rapalogues and CDK4/6 inhibitors have shown tolerability in the CNS, but have produced mixed results for treating CNS tumors.^{3,7,12,23}

Palbociclib has strong potential utility for treating DIPG, because RB disruption is uncommon in DIPG and cells typically lack functional p16^{INK4A}; key prerequisites for palbociclib efficacy. Furthermore, p16^{INK4A} has been shown to be repressed by aberrant histone H3.3K27M expression, which is a common mutation in DIPG.^{26,42} Rapalogues also have intrinsic potential for treating DIPG and can be used to overcome the characteristic hyperactivated PI3K/AKT/mTOR signaling exhibited in DIPG cells.³⁷ For example platelet-derived growth factor receptor (PDGFR) is commonly overexpressed in DIPG and drives cancer cell proliferation via the PI3K/AKT/mTOR signaling pathway. Combining rapalogues with CDK4/6 inhibitors has proven effective previously,^{4,18} and in accord with other groups, we found that palbociclib potentiates the effects of temsirolimus and may facilitate circumvention of its downstream repression by AKT. We propose palbociclib negates intrinsic CDK4-cycD1 complex inhibition of tuberous sclerosis complex 2,⁴¹ thus increasing mTORC1 inhibition and sensitization to temsirolimus. Furthermore, the decreased cycD1 and CDK4/6

expression, along with augmented hypophosphorylation of RB observed following our combination treatments, would serve to decrease mTOR activity further within this paradigm.

In addition to palbociclib potentially supplementing the effects of temsirolimus, temsirolimus may also potentiate palbociclib. Documented undesirable effects that have curtailed interest in palbociclib as a cancer treatment, such as attenuation of its own antimitogenic effects through increased cellular metabolism, can be abrogated by mTOR inhibition.²⁴ This was observed in our experiments via enhanced RB activation. We hypothesize this may arise through the capacity of temsirolimus to block global pro-mitogenic cellular effects caused by palbociclib (as described above), in addition to a blockade of cycD1 expression, which is positively-regulated by mTOR and would hence be restricted by temsirolimus. With a cycD1 deficit, there would be a propensity for further restriction of CDK4/6-cycD1 complex formation and hence decreased RB phosphorylation. Interestingly, if CDK4/6-cycD1 complexes form part of the mTOR regulation machinery in DIPG cells, and palbociclib and temsirolimus synergize to drive a significant deficit of cycD1, then ultimately a negative feedback loop would ensue, whereby decreased cycD1 expression intensifies mTOR inhibition, and mTOR inhibition decreases cycD1 expression. Thus, palbociclib and temsirolimus potentially elicit an antimitogenic positive feedback loop when used concurrently, with palbociclib inducing rapalogue sensitization, and temsirolimus inhibiting the negative side-effects of palbociclib. We propose that pathway cross-regulation and mutual potentiation of the inhibition of target proteins, drives increased antiproliferative effects and the observed shift towards cytotoxicity in cancer cells.

Despite advocating that palbociclib potentiates the inhibitory effects of temsirolimus on mTOR and facilitates circumvention of AKT-driven restoration of mTORC1 activity, we also observed that concurrent use of palbociclib and temsirolimus increased activation of the mTORC2-associated rictor protein. This did not however appear to affect cell viability. This may in part be due to the relatively short 24-hour exposure time for most of our drug treatments, chosen principally to mimic patient drug infusion times, but which also illustrated rapid drug efficacy. Furthermore, where longer treatment times were used (i.e. Fig. 6 and 7), pro-mitogenic mTORC2 effects did not appear significant and cell proliferation continued to decline. We are unable to fully rationalize this discrepancy and intend further investigation, but suggest temsirolimus may directly inhibit mTORC2 in DIPG cells. Rapalogues have been found to have an inhibitory effect on mTORC2 in some cancer cells, dependent on high

expression of FKBP12.⁴³ Pertinently, FKBP12 is significantly overexpressed in childhood astrocytomas.⁴⁴

Conclusion

Taken together, our results support the notion that the best clinical potential for developing an efficacious DIPG therapeutic will stem from concurrent inhibition of multiple oncogenic pathways. Our data suggest palbociclib and temsirolimus synergistically reduce cell viability in SF7761, SF8628 and SU-DIPG IV cells primarily through increased cytostatic effects that drive down cell proliferation, with some cytotoxicity likely stemming from the loss of metabolic compensatory pathways. As this project is principally an *in vitro* study, further investigation is required to translate these findings to the clinical setting. Frequently, treatment failures occur because agents cannot cross the blood–brain barrier due to efflux transporters. The ability of palbociclib to sufficiently enter the brain remains a concern,²⁶ but mTOR inhibitors have been shown to ameliorate palbociclib efflux and facilitate increased drug concentrations in the CNS.^{18,45} Nevertheless, our results demonstrate that simultaneous inhibition of the CDK4/6-cycD1-RB and PI3K-AKT-mTOR pathways is a feasible approach to DIPG treatment.

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Disclosure

S.S. Gill is a consultant for Renishaw PLC. N.U. Barua is a consultant for Renishaw PLC. No potential conflicts of interest were disclosed by the other authors. This study was funded by Abbie's Army DIPG brain tumor research charity (to NU Barua and AS Bienemann), the Functional Neurosurgery Research Fund held by Southmead Hospital Charities (SS Gill), the Gatsby foundation (to WGB Singleton) and joint funding from the Medical Research Council and

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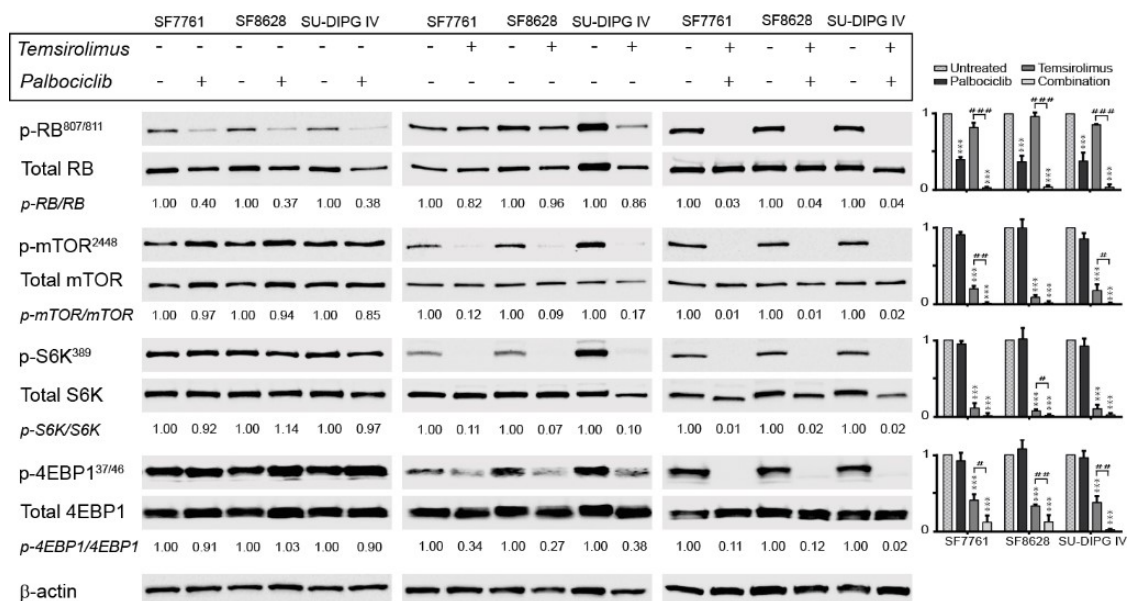


Figure 1 CDK4/6 and mTOR inhibitors cross-regulate, leading to enhanced inhibition of the respective pathways when used in combination.

Notes: SF7761, SF8628 and SU-DIPG IV cells were treated with either vehicle, 0.5 μ M palbociclib, 10 μ M temsirolimus or coadministered of 0.5 μ M palbociclib with 10 μ M temsirolimus for 24 hours prior to total protein extraction and immunoblot analyses. Quantitative densitometry values are shown below each blot. Densitometry graphs compare single-agent treatments and combination treatment to vehicle treated controls (right). β -actin was used as a loading control. Data are the mean \pm SEM of triplicate determinations in all instances. *, $p < 0.05$; **, $p < 0.01$; ***, $p < 0.001$ (compared to respective vehicle treatment). #, $p < 0.05$; ##, $p < 0.01$; ###, $p < 0.001$ (compared to respective temsirolimus single-agent treatment).

Abbreviations: RB, retinoblastoma protein; CDK, cyclin-dependent kinase; mTOR, mechanistic target of rapamycin protein; S6K, p70 ribosomal protein S6 kinase; 4EBP1, eukaryotic translation initiation factor 4E-binding protein 1; rictor, rapamycin-insensitive companion of mammalian target of rapamycin; p-, phosphorylated form of protein.

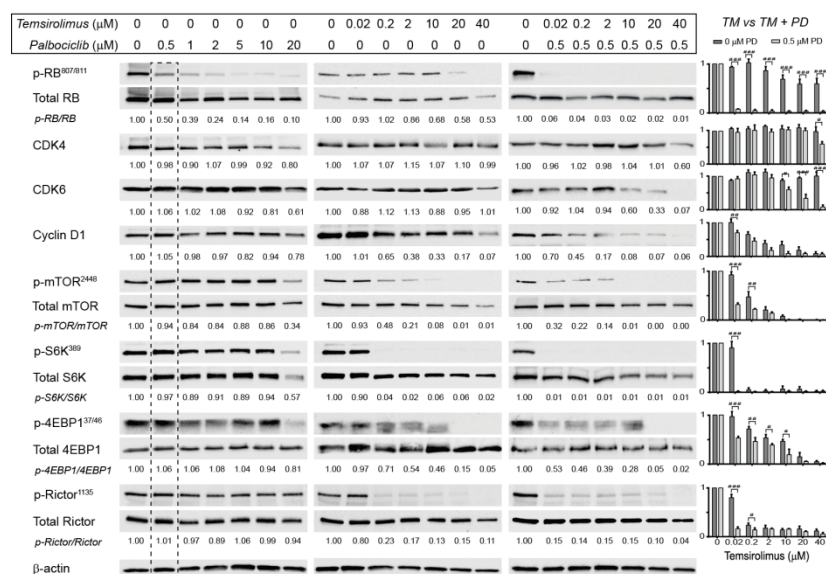


Figure 2 Coadministration of palbociclib with temsirolimus yields beneficial enhancement of CDK4/6 and mTOR inhibition in DIPG cells.

Notes: SF7761 cells were treated with a range of doses of palbociclib, temsirolimus or temsirolimus coadministered with 0.5 μ M palbociclib for 24 hours prior to total protein extraction. Immunoblot analyses showed palbociclib and temsirolimus effectively inhibited CDK4/6-cycD1-RB and mTOR signaling pathways in DIPG cells, respectively, while each drug also exhibits inherent attributes to disrupt the other's target protein, leading to an ostensible synergistic outcome when used in combination. Quantitative densitometry values are shown below each blot. Densitometry graphs compare single-agent temsirolimus treatment and temsirolimus given in combination with 0.5 μ M palbociclib (right). Dotted line emphasizes 0.5 μ M palbociclib single-agent results versus combination effects. β -actin was used as a loading control. Data are the mean \pm SEM of triplicate determinations in all cases. #, $p < 0.05$; ##, $p < 0.01$; ###, $p < 0.001$ (compared to respective temsirolimus single-agent treatment).

Abbreviations: RB, retinoblastoma protein; CDK, cyclin-dependent kinase; mTOR, mechanistic target of rapamycin protein; S6K, p70 ribosomal protein S6 kinase; 4EBP1, eukaryotic translation initiation factor 4E-binding protein 1; rictor, rapamycin-insensitive companion of mammalian target of rapamycin; p-, phosphorylated form of protein. PD, palbociclib; TM, temsirolimus.

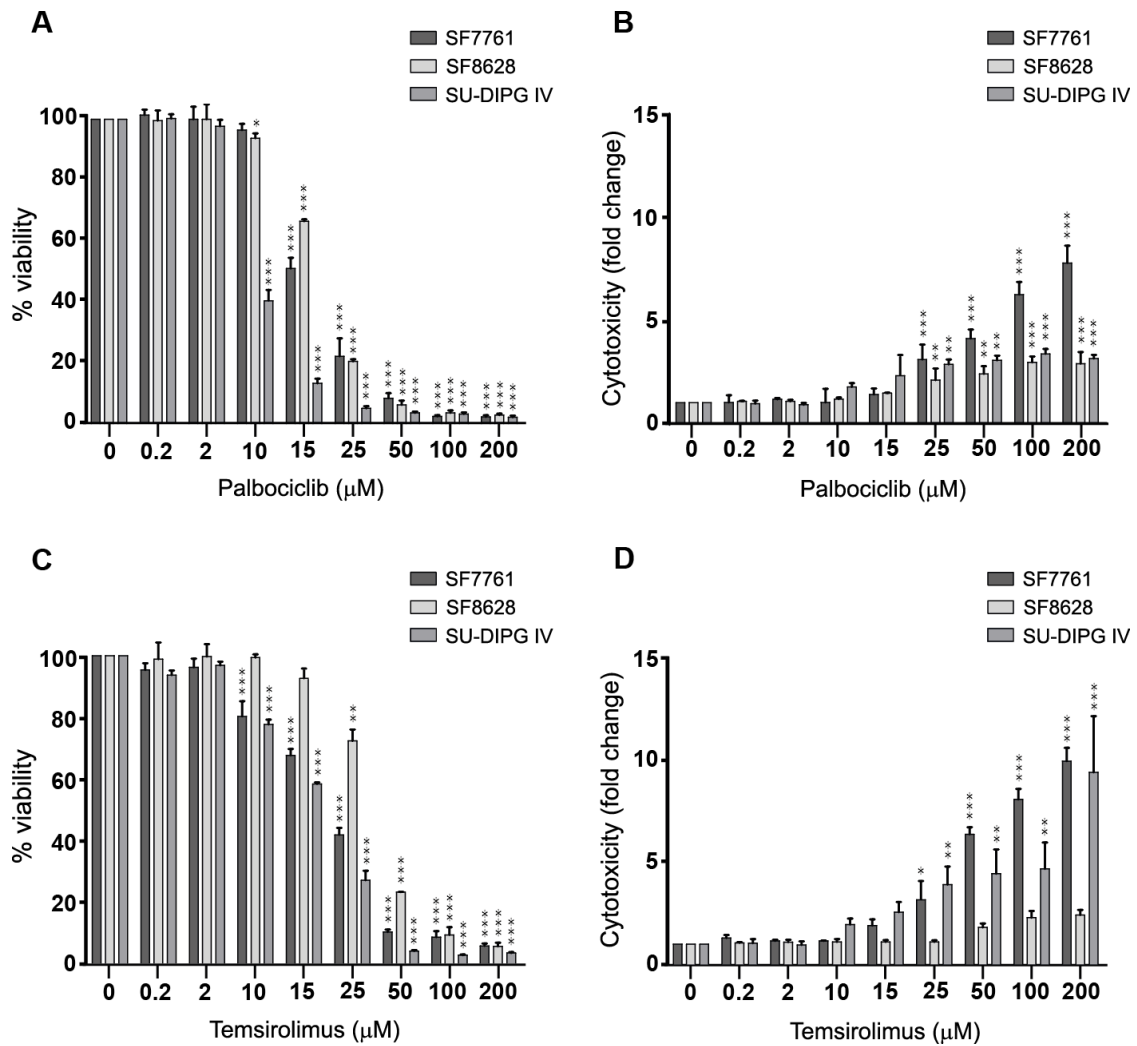


Figure 3 Palbociclib and temsirolimus treatments as monotherapies reduce cell viability in DIPG.

Notes: A, calcein-AM cell viability assays were performed on SF7761, SF8628 and SU-DIPG-IV DIPG cell lines treated with increasing concentrations of palbociclib for 24 hours. B, drug cytotoxicity was measured following 24 hours treatment with palbociclib, using ethidium homodimer-1 (ethD-1). C, cell viability assays were performed on SF7761, SF8628 and SU-DIPG IV cell lines treated with increasing concentrations of temsirolimus for 24 hours. D, drug cytotoxicity was measured following 24 hours treatment with temsirolimus, using ethD-1. Data are the mean \pm SEM of triplicate determinations. *, $p < 0.05$; **, $p < 0.01$; and ***, $p < 0.001$ (compared to the respective vehicle treatment).

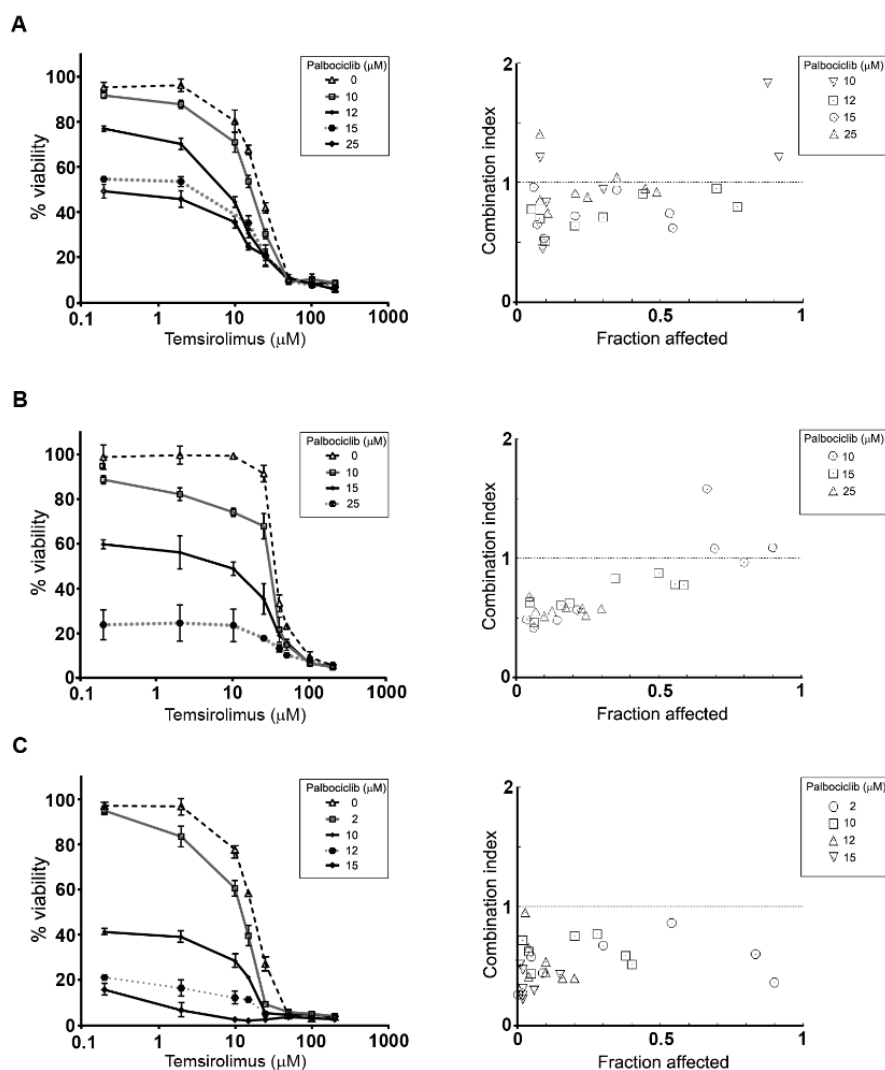


Figure 4 Coadministration of palbociclib and temsirolimus decreases cell viability synergistically.

Notes: SF7761 (A), SF8628 (B) and SU-DIPG IV (C) cells were treated with single-agent temsirolimus at increasing concentrations (0.2-200 μM) or the same concentration range combined with a single fixed dose of palbociclib (either 2, 10, 12, 15 or 25 μM). Cell viability was assessed using calcein-AM staining. A combination index score was assigned to each of the different combinations and is shown on the right. Data are the mean \pm SEM of triplicate determinations.

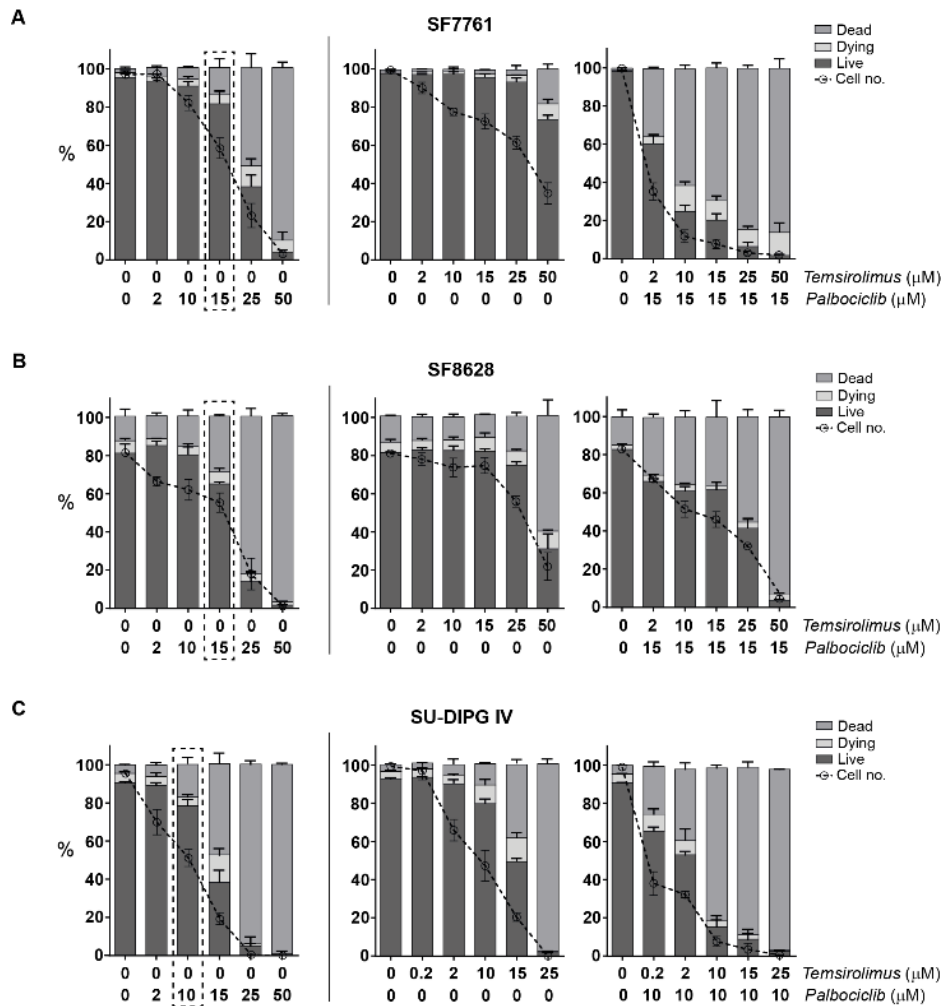


Figure 5 Coadministration of palbociclib and temsirolimus decreases cell viability principally through restricting cell proliferation.

Notes: Bivariate flow cytometric analyses of cell viability in SF7761 (A), SF8628 (B) and SU-DIPG IV (C) cells. Cells were treated with palbociclib (0–50 mM), temsirolimus (0–50 mM) or both (temsirolimus 0–50 mM, with a fixed dose of either 10 mM or 15 mM palbociclib, dependent on cell line sensitivity). Analysis of “% live cells” (calcein-AM positive), “% dead cells” (ethD-1 positive) and “% dying cells” (double positive) was performed. The dotted line box in each of the panels on the left-hand side emphasizes the dose of palbociclib used for the combinatorial analyses. Data are the mean \pm SEM of triplicate determinations.

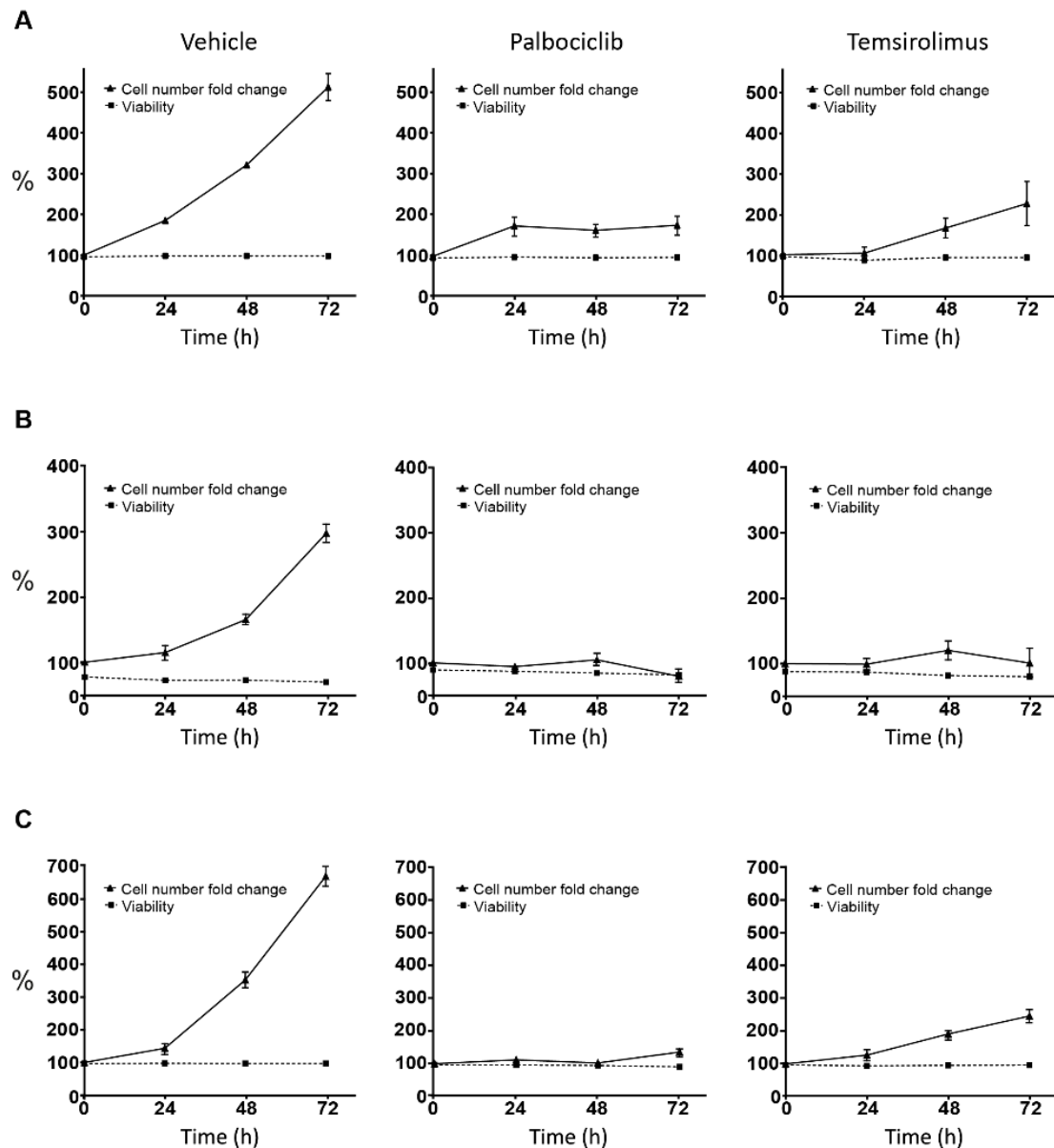


Figure 6 Palbociclib and temsirolimus inhibit the growth of DIPG cells cytostatically.

Notes: SF7761 (A), SF8628 (B) and SU-DIPG IV (C) cells were treated with vehicle, palbociclib (2 μ M) or temsirolimus (10 μ M for SF7761 and SF8628; 0.2 μ M for SU-DIPG IV) for 0-72 hours. Cells were then counted and stained with calcein-AM to deduce % viability, via flow cytometry. Data are the mean \pm SEM of triplicate determinations in all cases.

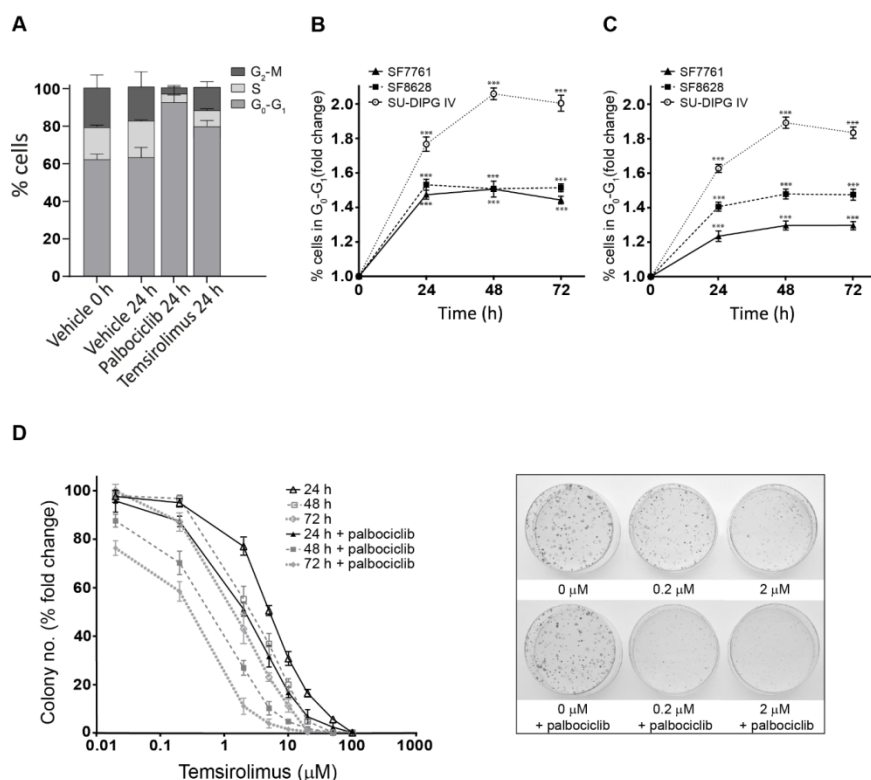


Figure 7 Palbociclib and temsirolimus treatments lead to cell cycle arrest.

Notes: A, DRAQ5 fluorescent dye was used to perform flow cytometric cell cycle analysis on SF7761 cells treated with either vehicle, 2 μM palbociclib or 10 μM temsirolimus for 24 hours. B, DRAQ5 cell cycle analysis of SF7761, SF8628 and SU-DIPG IV cells treated with 2 μM palbociclib demonstrated a significant increase in the percentage of cells in the G₀-G₁ phase over time. C, DRAQ5 cell cycle analysis of SF7761 and SF8628 cells treated with 10 μM temsirolimus, and SU-DIPG IV cells treated with 0.2 μM temsirolimus demonstrated a significant increase in the percentage of cells in the G₀-G₁ phase over time. D, SU-DIPG IV cells were treated with temsirolimus, or the combination of temsirolimus and 0.5 μM palbociclib for 24-72 hours, and colonies counted after 14 days. Representative plates following 72 hours treatment are shown (right). Data are the mean ± SEM of triplicate determinations in all cases. *, p < 0.05; **, p < 0.01; and ***, p < 0.001 (compared to respective vehicle treatment).

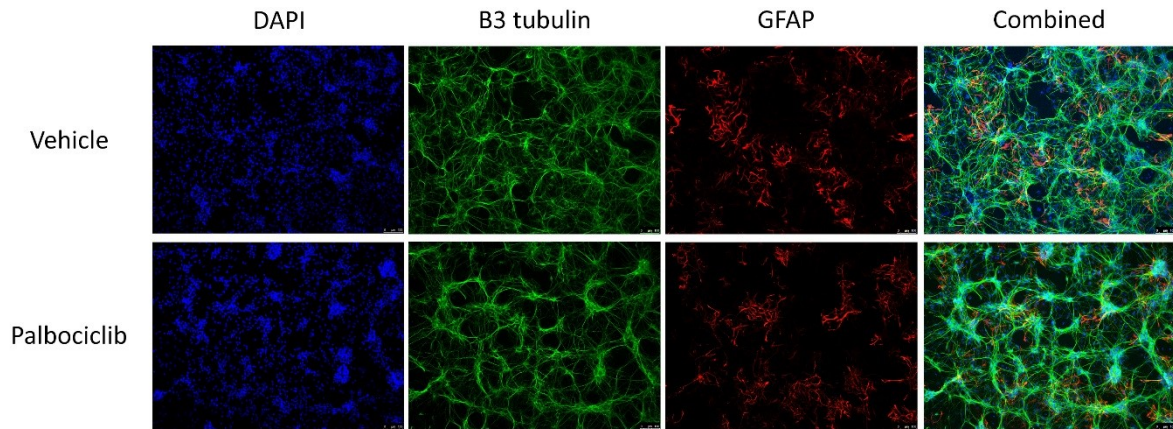


Figure 8 Treatment of rat hippocampal primary cultures with palbociclib did not elicit cytotoxicity.

Notes: Primary hippocampal culture treated with vehicle solution or 2 μ M palbociclib for 24 hours. Intact neuronal networks and normal glial cell morphology was observed in cultures treated with palbociclib, suggesting no significant toxicity. Cell nuclei were stained with DAPI (blue); neurons were stained with B3 tubulin antibody (green); glial cells were stained with GFAP antibody (red/orange).

Abbreviations: DAPI, 4',6-diamidino-2-phenylindole; B3 tubulin, tubulin, beta III isoform; GFAP, glial fibrillary acidic protein.

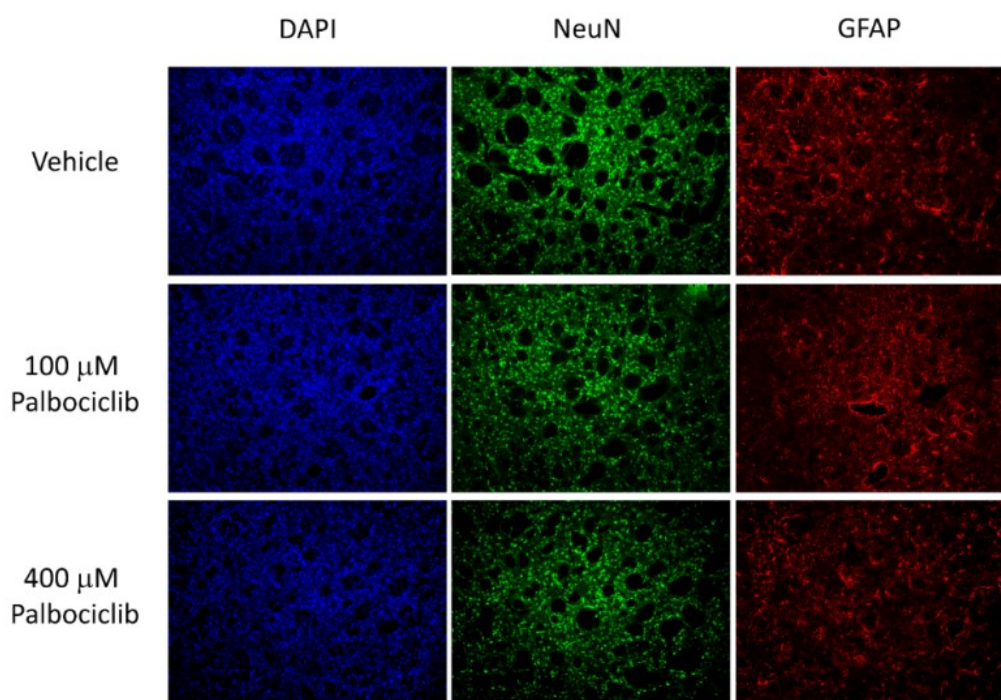


Figure 9 Direct delivery of palbociclib into rat brain did not elicit discernible toxicity.

Notes: Palbociclib was infused into the striatum of rat brains at a rate of 1 μ l / min (5 μ l total) at either 100 μ M or 400 μ M. Tissue was harvested after 48 hours and analyzed using immunofluorescence staining. Tissue morphology appeared normal in rats dosed with palbociclib, suggesting no significant toxicity. Cell nuclei were stained with DAPI (blue); neurons were stained with NeuN antibody (green); glial cells were stained with GFAP antibody (red).

Abbreviations: DAPI, 4',6-diamidino-2-phenylindole; NeuN, neuronal nuclear protein; GFAP, glial fibrillary acidic protein.

Table 1 IC₅₀ values for 24 hours single-agent and combination treatments in SF7761, SF8628 and SU-DIPG IV cells

Drug treatment	IC ₅₀ (μM)	IC ₅₀ (μM)	IC ₅₀ (μM)
	SF7761	SF8628	SU-DIPG IV
Palbociclib (0.2-200 μM)	16.55	16.50	8.77
Temsirolimus (0.2-200 μM)	20.88	36.62	16.83
Temsirolimus (0.2-200 μM) + 2 μM Palbociclib	-	-	11.52
Temsirolimus (0.2-200 μM) + 10 μM Palbociclib	15.83	24.95	0.16
Temsirolimus (0.2-200 μM) + 12 μM Palbociclib	4.82	-	0.002
Temsirolimus (0.2-200 μM) + 15 μM Palbociclib	0.89	2.03	0.002
Temsirolimus (0.2-200 μM) + 20 μM Palbociclib	-	0.001	-
Temsirolimus (0.2-200 μM) + 25 μM Palbociclib	0.44	-	-